

PRINTER RUSH

(PTO ASSISTANCE)

2nd request

Application : 09/945396

Examiner : Le

GAU : 1641

From: MR

Location: IDC FMF FDC

Date: 01/12/05

Tracking #: 06062095 Week Date: 01-03-05

DOC CODE	DOC DATE	MISCELLANEOUS
<input type="checkbox"/> 1449	_____	<input type="checkbox"/> Continuing Data
<input type="checkbox"/> IDS	_____	<input type="checkbox"/> Foreign Priority
<input type="checkbox"/> CLM	_____	<input type="checkbox"/> Document Legibility
<input type="checkbox"/> IIFW	_____	<input type="checkbox"/> Fees
<input type="checkbox"/> SRFW	_____	<input checked="" type="checkbox"/> Other Seq. listing
<input type="checkbox"/> DRW	_____	
<input type="checkbox"/> OATH	_____	
<input type="checkbox"/> 312	_____	
<input type="checkbox"/> SPEC	_____	

[RUSH] MESSAGE: Please provide Sequence listing pages to be used for publication. CRPL dated 11-21-02 after SPEC dated 11-13-02 contains "Raw Sequence listing."

Thank you,
MR

[XRUSH] RESPONSE: _____

/M/R

INITIALS: X

NOTE: This form will be included as part of the official USPTO record, with the Response document coded as XRUSH.

REV 10/04

Log 10/12/05

10/12/05

REMARKS

A CRF and paper version of the "Sequence Listing" are provided herewith. Applicants hereby state that the contents of the CRF and the paper version of the CRF are identical. The sequences are plainly present in the original Specification. No new matter has been added.

The present CFR CD was prepared and validated by Patentin version 3.1.1.6.

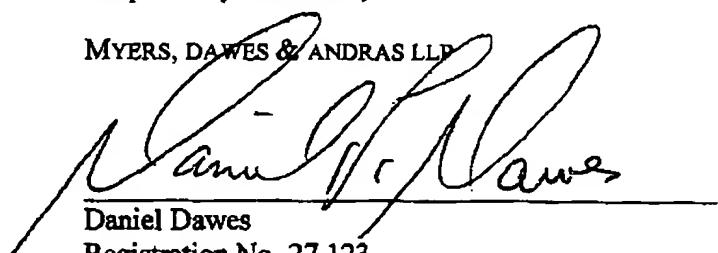
The specification has also been amended to insert "sequence ID NO:" indicators no other changes have been made, no new matter has been added.

Applicant respectfully submits that the case is now in condition for allowance and requests an early notification of the same. Questions, suggestions, and comments from the Examiner are welcomed. If the Examiner believes that a telephone conference would help further the prosecution of the case, the Examiner is requested to contact the undersigned attorney at the listed telephone number.

November 4, 2002

Respectfully submitted,

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(Date of Deposit)

Nancy V. Mr. Elrath
Nancy V. Mr. Elrath

Date of Signature 11-4-02

Following is the marked-up version of the amended specification paragraph.

Fig. 7 is a graph of phosphorylation as a function of time showing the separation of three different specific kinase reporter substrates from a sample of the contents of a *Xenopus laevis* oocyte 46' that had been previously microinjected to contain ~1~~2~~M Fl-sPKC, ~330 nM Fl-sPKA, and ~10 nM Fl-scdc2K. Fl-sPKA, a specific reporter for protein kinase A (PKA) activity has the sequence FL-Lys-Arg-Arg-Glu-Ile-Leu-Ser-Arg-Arg-Pro-Ser-Tyr-Arg (sequence ID NO:1), and was derived from the CREB protein. Fl-scdc2K, a specific reporter for cdc2 kinase (originally identified genetically as cell division cycle mutant 2) has the sequence Fl-Gly-Gly-Gly-Arg-Ser-Pro-Gly-Arg-Arg-Arg-Lys (sequence ID NO:2), and comprises a consensus phosphorylation site derived from several proteins. The underlined serine residues are the sites of phosphorylation. The peptides were synthesized and labeled with fluorescein as described for Fl-sPKC, except that Fl-scdc2K was labeled with the mixed 5- and 6-isomers of carboxyfluorescein succinimidyl ester (100-200 mg/ml, Molecular Probes, Eugene OR); thus, Fl-scdc2K consisted of two isomeric forms. A peak 162 and a peak 172 were identified by their migration times as observed when injected into oocytes 46' singly (not shown). The first doublet, peaks 162 and 164, corresponds to two isomers of either phosphorylated or nonphosphorylated Fl-scdc2K. The second doublet, peaks 166 and 168, corresponds to two isomers of the other form of Fl-scdc2K. One peak 170 represents nonphosphorylated Fl-sPKC, and one peak 172 represents nonphosphorylated Fl-sPKA.